

AperTO - Archivio Istituzionale Open Access dell'Università di Torino

Does Adsorption at Hydroxyapatite Surfaces Induce Peptide Folding? Insights from Large-Scale B3LYP Calculations

This is the author's manuscript

Original Citation:

Availability:

This version is available <http://hdl.handle.net/2318/106459> since 2016-08-08T09:35:40Z

Published version:

DOI:10.1021/ja302262y

Terms of use:

Open Access

Anyone can freely access the full text of works made available as "Open Access". Works made available under a Creative Commons license can be used according to the terms and conditions of said license. Use of all other works requires consent of the right holder (author or publisher) if not exempted from copyright protection by the applicable law.

(Article begins on next page)



UNIVERSITÀ DEGLI STUDI DI TORINO

This is an author version of the contribution published on:

Questa è la versione dell'autore dell'opera:

Rimola, A.; Aschi, M.; Orlando, R.; Ugliengo, P. Does Adsorption at Hydroxyapatite Surfaces Induce Peptide Folding? Insights from Large-Scale B3LYP Calculations. *Journal of the American Chemical Society* **2012**, *134*, 10899-10910. 10.1021/ja302262y

The definitive version is available at:

La versione definitiva è disponibile alla URL:

<http://pubs.acs.org/doi/abs/10.1021/ja302262y?prevSearch=%255BContrib%253A%2Bugliengo%255D&searchHistoryKey=>

Does Adsorption at Hydroxyapatite Surfaces Induce Peptide Folding?

Insights from large scale B3LYP calculations

Albert Rimola^{1,2}, Massimiliano Aschi³, Roberto Orlando² and Piero Ugliengo^{*2}

¹*Departament de Química, Universitat Autònoma de Barcelona, 08193 Bellaterra, Spain.*

²*Dipartimento di Chimica, NIS Centre of Excellence and INSTM (Materials Science and Technology) National Consortium, University of Torino, Via P. Giuria 7, 10125 Torino, Italy.*

³*Department of Chemistry, Chemical Engineering and Materials, University of L'Aquila, Via Vetoio (Coppito 1), 67010 L'Aquila, Italy.*

Abstract

Large scale periodic quantum mechanical calculations (509 atoms, 7852 atomic orbitals) based on the hybrid B3LYP functional focused on the peptide folding induced by the adsorption on the (001) and (010) hydroxyapatite (HA) surfaces give interesting insights on the role of specific interactions between surface sites and the peptide, which stabilize the helix conformation over the “native” random coil ones for *in silico* designed model peptides. The two peptides were derived from the 12-Gly oligomer, with one (P1: C-tGGKGGGGGGEGGN-t) and two (P2: C-tGGKGGKEGGEGGN-t) glutamic acid (E) and lysine (K) residue mutations. The most stable gas-phase “native” conformation for both peptides resulted in a random coil (RC) structure, with the helix (H) conformation being $\approx 100 \text{ kJ mol}^{-1}$ higher in free energy. The two peptide conformations interact with the HA (001) and (010) surfaces by C=O groups *via* Ca^{2+} ions, by hydrogen bond between NH_2 groups and the basic PO_4^{3-} groups and by a relevant fraction due to dispersion forces. Peptide adsorption was studied on the dry (001) surface, the wet one envisaging 2 H_2O *per* surface Ca^{2+} and, on the latter, also considering the adsorption of micro-solvated peptides with 4 H_2O molecules located at sites responsible of the interaction with the surface. The P1 mutant does prefer to be adsorbed as a random coil by $\approx 160 \text{ kJ/mol}$, whereas the reverse is computed for P2, preferring the helix conformation by $\approx 50 \text{ kJ/mol}$. Adsorption as helix of both P1 and P2 mutants brings about proton transfer towards the HA surfaces with a large charge transfer component to the interaction energy.

Introduction

The biological activity of a given protein is partly due to its secondary structure and conformational state. Peptide chains are rather flexible so that the interest in finding novel ways that force the protein folding in a well-defined state has increased considerably. To be folded a protein needs extra-interactions that compensate for the entropy loss associated with the folding of the polypeptide chain. Among the most common adopted constraint techniques, such as metal-ion complexation¹⁻⁵ or S-S bonds,⁶⁻⁸ the interaction of proteins with inorganic surfaces has demonstrated to be a fruitful strategy to stabilize selected folded states.⁹⁻¹⁹ This, nevertheless, remains a challenging goal since the protein-surface binding usually leads to denaturation and the loss of the biological functionality. Due to that, de novo peptide design techniques has emerged as a useful approach to assemble peptides with a defined structure and orientation on the surface,²⁰ as the synthetic peptide may contain, at very strategic positions, amino acidic residues that when in contact with the surface establish extra intermolecular interactions that drive and stabilize the folded protein.

The surface-induced peptide folding is a promising tool to: i) obtain peptide monolayers exhibiting free-interacting sequences that may display specific biochemical recognition phenomena; ii) induce a specific bioactivity of the protein which is missing in its unfolded state. Both issues have several important applications in biosensing, biotechnology, biocatalysis, biomaterials, tissue engineering, nanotechnology and proteomics. Interestingly, the peptide folding induced by mineral surfaces and the subsequent activation of a potentially “hidden” bioactivity, might have triggered the first biocatalytic reactions in a primordial Earth (in absence of life), hence giving rise to the emergence of the metabolic cycles, a crucial aspect for the origin of life.^{21,22}

Several experimental studies performed with high-resolution spectroscopic techniques such as circular dichroism (CD) and solid state nuclear magnetic resonance (ssNMR) have been reported in the literature. They were focused on the formation of peptide secondary structures induced by the presence of surfaces and observed the formation of helical structures from synthetic peptides when interacting with hydrophobic surfaces⁹ and cavities,¹⁰ silica nanoparticles,^{11,12} functionalized gold nanoparticles^{13,14} and hydroxyapatite.¹⁵ Such conformational changes have also been observed in natural proteins adsorbed on gold nanoparticles¹⁶ and hydroxyapatite.¹⁷⁻¹⁹ Despite the novelty of the works and the useful information provided, detailed atomistic features related to the binding mechanisms and the anchoring points between peptides and the surfaces are very difficult to attain. One way to provide this missing information is by classical molecular dynamics-based studies,²³⁻²⁶ focusing on the dynamic behavior of proteins in water and when docked to the surfaces. However, the adoption of classical force fields (FF), needed to handle the large number of atoms involved, inevitably lower the accuracy of the results, as actual FF cannot cope with bond breaking/making which may indeed occur between side-chain functionalities and the surfaces. One exception is by adopting a reactive force field like ReaxFF²⁷ which is able, by construction, to handle bond rupture/formation in a way mimicking quantum mechanical (QM) calculations. Indeed, very recently, Monti and co-workers²⁸ have extended ReaxFF to study the interaction of glycine and diglycine on the TiO₂ rutile (110) face using a newly ReaxFF force field for glycine.²⁹ QM calculations are the obvious choice to keep almost constant accuracy in the diverse situations occurring by these complex simulations, at the price of a very high computational cost.^{25,26} For these reasons we have adopted for this work CRYSTAL09 in its massive parallel version which is able to exploit hundreds of cores to solve the Schrödinger equation for periodic systems.³⁰

We have been inspired by the work of Cappriotti et al.,¹⁵ in which it was shown that a rich- γ -carboxyglutamic synthetic peptide assumes a random coil structure in solution whereas it becomes reorganized into an α -helix when in contact with HA surfaces. Our target was to study such a process entirely *in silico*, by adopting a simplified version of the peptides and focusing on the two most common HA surfaces, namely the (001) and the (010) ones, respectively. We chose the hybrid functional B3LYP³¹ in a periodic context, adopting Gaussian basis sets of sufficiently high quality. Due to the complexity of the system and of high computational cost we had to compromise on both the solvation state of the entire system and to limit the calculation to static optimization, as *ab initio* MD would have been too computational demanding. Our strategy was, first, to study the surface induced folding process entirely in gas-phase and then to micro-solvate the system with 8 water molecules at the sites responsible of the interactions for the gas-phase processes. In this way, the key role of the first solvation shell in crucial sites has been addressed. We are aware that this approach has serious limitations due to the above simplifications; notwithstanding, as it will be shown in the paper, a number of important energetic and structural details have been characterized.

Methods

In the following only essential computational details are provided, a more extensive description being provided in the Supplementary Information (SI). All the periodic calculations for the interaction of the considered peptides with the HA (001) and (010) surfaces have been performed with the massive parallel version of the *ab initio* code CRYSTAL09,^{32,33} which solves the Schrödinger equation for molecules, polymers, slabs and bulks using Gaussian basis sets. All the SCF calculations and geometry optimizations were performed with the B3LYP density functional method.^{31,34}

Geometry optimizations were performed using a Gaussian basis set of polarized double- ζ quality already adopted in previous studies.³⁵ Additionally, to increase the accuracy in computing the adsorption energies, single-point energy calculations on the optimized structures using the Ahlrichs and coworkers VTZP basis set³⁶ were carried out which, in addition, significantly reduces the BSSE as we demonstrated recently.³⁷ With the VTZP basis set the number of atomic orbitals was 7852 which required distributing the Fock diagonalization over many cores. Geometries were optimized by relaxing the internal coordinates within PI symmetry keeping the lattice parameters fixed at the values of the optimized bare surfaces.

In a periodic treatment, the adsorption energy ΔE_{ads} per unit cell per adsorbate is defined as:

$$\Delta E_{\text{ads}} = E(\text{SP//SP}) - E(\text{S//S}) - E_{\text{m}}(\text{P//P})$$

where $E(\text{S//S})$ is the energy of a bare HA slab S in its optimized geometry, $E_{\text{m}}(\text{P//P})$ is the molecular energy of a free peptide P in its optimized geometry and $E(\text{SP//SP})$ is the energy of the considered HA/P system in its optimized geometry (the symbol following the double slash identifies the optimized geometry at which the energy has been computed). ΔE_{ads} is a negative quantity for a bound system and can be recast in terms of deformation cost of the surface, the adsorbate, the lateral adsorbate-adsorbate interactions, and the interaction between the predeformed constituents. The final expression $\Delta E_{\text{ads}}^{\text{C}}$, inclusive of the BSSE correction reads:

$$\Delta E_{\text{ads}}^{\text{C}} = \Delta E_{\text{ads}} + \text{BSSE}$$

It is worth mentioning that for periodic systems the BSSE can only be evaluated for cases in which both the adsorbate and the surface remains electro neutral, so that all processes in which bond making/breaking occurs are excluded. In this work, only adsorption of peptides in the random coil forms remain neutral when adsorbed (no

proton transfer) and, accordingly, only for these cases the BSSE have been computed. For the helix conformations, on the contrary, proton transfers do indeed occur preventing the BSSE evaluation. Notwithstanding, to keep balance between interaction energies, for the helix cases the highest percentage of BSSE computed for the corresponding random coil cases was used to estimate the final $\Delta E_{\text{ads}}^{\text{C}}$ values.

B3LYP, as almost all standard hybrid and pure gradient corrected functionals, cannot cope with the dispersion energy component to the interaction energy, so that the empirical correction ΔD suggested by Grimme³⁸ and re-parameterized by some of us for periodic systems,³⁹ has been added to the BSSE-corrected adsorption energies $\Delta E_{\text{ads}}^{\text{C}}$ in a posteriori fashion to obtain the final $\Delta E_{\text{ads}}^{\text{C}} + \Delta D$. In this way, the dispersive contribution is inserted in a non self-consistent way to the adsorption energy, which, somehow, underestimates its relevance (see also SI for more details).

Conformational study of peptide in the gas-phase at room temperature was performed using MD simulations with the Gromos 53a6 force-field⁴⁰ and the Gromacs package.⁴¹ The peptide, initially folded and minimized in the helix conformation, was simulated for 160.0 ns with a time step of 2.0 fs at a temperature of 300.0 K kept constant with the Berendsen thermostat⁴² with a time constant equal to the integration step. The LINCS algorithm⁴³ was used to constrain all the bonds. Conformational analysis was carried out using Essential Dynamics⁴⁴ by projecting the whole trajectory onto the subspace (essential space) constructed using the subset of eigenvectors, obtained by the all atoms covariance matrix diagonalization, characterized by the highest eigenvalues. This operation produces different basins of probability approximately corresponding to well-defined tridimensional peptide structures. Hence, the random-coil structure utilized in this work represented the most populated

probability (lowest free energy) basin. Further details are provided in the supporting information.

Results

In silico designed peptides

The (001) and (010) HA surfaces are the most relevant ones from a biological viewpoint and have been chosen here as surface models to study the folding induced by the adsorption of the model peptides. HA has been chosen as a test system for two reasons: i) it is the natural major inorganic constituent of bone and teeth and, accordingly, the natural choice to study biomolecule/biomaterial interactions; ii) the crystalline structure of HA ($\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2$) exhibits calcium ions periodically spaced at a distance close to the intervals of the repeating turns in an α -helix polypeptide. For point ii), it is expected that sequences that contain residue side chains with a large Ca^{2+} affinity will interact favorably with the HA surfaces. Additionally, HA also contains phosphate groups, so that peptide residues containing groups behaving as H-bonding donors strongly interact with the HA surfaces. In summary, amino acids containing either acidic or basic residues are expected to have high affinity for the HA surfaces.

Indeed, calculations in both gas-phase and under micro-solvation conditions (few water molecules) carried out by some of us show that single amino acids either acidic (namely, aspartic and glutamic acids) or basic (namely, lysine and arginine) interact strongly with HA surfaces.^{35,45-48} This is in agreement with experimental measurements indicating that electrostatic complementarity between protein and the surface fields is the leading force that drive the adsorbate to the surface and, eventually, causing the protein to fold as a helix.¹¹⁻¹⁵

To study a possible role of the HA surfaces on the protein folding, we designed an *in silico* Gly polymer in its α -helix conformation, as shown in Figure 1. We firstly optimized the geometry of the surface-free polyglycine whose unit cell contains 7 independent Gly residues (linear group, *P1*). The resulting optimized structure (Figure 1, H-[G]_p structure), is indeed a perfect α -helix with the typical $i+4 \rightarrow i$ H-bonds feature. The infinite helix was then cut to define a small peptide with 12-Gly residues (structure H-[G]_m, Figure 1), showing three helical turns. The dangling bonds resulting from the polymer to the 12-mer transformation have been healed by H (NH₂ terminus) and OH (COOH terminus), labeled by stars in Figure 1. The choice of 12-mer is consistent with the size of the HA cell chosen to represent the (001) and (010) surfaces (12-mer length of 22 Å vs 24 Å for the diagonal of the HA surface cell). The 12-Gly polypeptide is highly hydrophobic, so in order to provide residues apt to interact with the HA surfaces (*vide supra*) the peptide has been mutated by substituting some of the Gly with glutamic acid (E) and lysine (K). In one mutant (P1: C-tGGKGGGGGEGGN-t), Figure 1) only one E and K residues were added, while in the other (P2: C-tGGKGGKEGGEGGN-t), Figure 1) two E and K were included. In order to maximize the interactions with the HA surfaces, the E and K residues occupy positions in which their side chains remains on the same side of the main helix axis (see Figure 1).

Surface-free gas-phase helix folding

The optimized helix forms of P1 and P2 in gas-phase are shown in Figure 2a (H-P1) and Figure 2b (H-P2), respectively. They resulted from the optimization of the mutated peptides as cut forms of the periodic polyglycine. The structure of H-P1 has five consecutive $i + 3 \rightarrow i$ H-bonds and three consecutive $i + 4 \rightarrow i$ ones; structure H-P2 shows three consecutive $i + 3 \rightarrow i$ H-bonds and five consecutive $i + 4 \rightarrow i$ H-bonds.

Thus, peptides exhibit a mixture of 3_{10} and α -helix structures, as shown by the triangular and square shapes associated to the 3_{10} and α -helix structures, respectively (Figures 2c-f). Any attempt to convert the E and K residues in a zwitterionic state (by imposing KH^+ and E^- states) resulted in a spontaneous proton transfer back to restore the neutral K and E residues. This is in agreement with what is known about gas-phase instability of the zwitterionic form of free amino acids.⁴⁹

The method so far adopted forced the peptide to assume a helix conformation by construction. However, one has to establish whether this conformation is the most stable one for the free P1 and P2 peptides. An exhaustive conformational search by means of classical molecular dynamics (MD) simulations of both peptides have been run in gas-phase conditions to determine the most stable conformation. Results showed that both peptides prefer a random coil conformation (RC-P1 and RC-P2, Figure 2g and h, respectively) over the starting helix one. A Ramachandran plot of the ϕ and Ψ torsion angles for the four peptides (H-P1, H-P2, RC-P1, RC-P2) (Figure 2i) confirms that H-P1 and H-P2 values are within the region of the helix whereas RC-P1 and RC-P2 values are filling the whole region, showing character of random coil.

The random coil peptides obtained from the MD calculations have then been fully optimized at B3LYP level, so that the reaction energy ΔE_{rx} of RC-P1 \rightarrow H-P1 and RC-P2 \rightarrow H-P2 processes have been computed (see Table 1). QM data showed that the folding of both peptides from random coil to helix is endoenergetic ($\Delta E_{\text{rx}} > 0$). The addition of the dispersion contribution to the pure B3LYP energies (ΔD column, Table 1) further increases the endoenergetic character of the processes ($\Delta E_{\text{rx}} + \Delta D$ column, Table 1). The entropic contribution computed by standard statistical mechanics formulae using the harmonic vibrational frequencies should be considered as rather approximated as it does not include neither the anharmonicity character of many modes

characterized by large amplitude motion nor the configuration entropy due to conformations close in energy. Explorative calculations run with classical MD (see Figure S1, SI) revealed, however, that these factors are less important in gas-phase than in water. Within these approximations, the $(T \cdot \Delta S)$ resulted negative (about -20 kJ mol^{-1}), in agreement with the lower entropy content of the helix compared to the random coil. The final free reaction energies inclusive of the dispersive contribution are around 100 kJ mol^{-1} , irrespective of the mutants. Data in Table 1 also show the very good agreement between the reaction energy computed with cheap single point energy calculations at B3LYP/VTZP//B3LYP/double- ζ polarized basis set and those computed with the expensive fully optimized B3LYP/VTZP ones. Because of that, the subsequent energy evaluations have all been carried out as single point B3LYP/VTZP//B3LYP/double- ζ polarized basis set.

Structures and energetics of peptides adsorbed on dry HA surfaces

Previous results established, on a firm base, that the random coil conformation is more stable than the helix one for the considered gas-phase peptides. We have then studied whether the adsorption on the HA (001) and (010) surfaces may change the energetic of the considered reactions. Our previous experience^{35,45,46,48} indicates that the peptide/HA interactions are dictated partly by specific electrostatic interactions. For the helix forms, specific interactions of the K and E side chains with the Ca^{2+} and PO_4^{3-} surface groups are obviously the dominant ones. For the random coil peptides, due to the more compact structure, the most favorable sites of interactions with the surfaces are not self-evident. For these cases, we have computed the electrostatic potential maps (EPMs) of all the random coil peptides (Figure S2, SI) and used the electrostatic complementarity between the electric features of the peptide and that of the clean HA

surfaces to properly dock the peptides towards the surface. The present approach does not consider at all the obvious fact that the most stable conformation adopted by the peptide in gas-phase may be different from the one adopted on the HA surfaces. In essence, an exhaustive conformational search of the peptide on the HA would have been required in a more rigorous approach. Clearly this would have been extremely demanding even for classical force field calculations and impossible to perform with our approach based on static minimization calculations.

The resulting structures for the RC-P1 and RC-P2 forms interacting with the HA (001) and (010) surfaces are shown in Figures 3 and 4, respectively. The interactions between the peptides and the HA surfaces are through two C=O groups belonging to the backbone chain of the peptides and Ca²⁺ ions of the surfaces. These carbonyl groups are those exhibiting the deepest negative electrostatic potentials in the EPMs (Figure S2, SI). H-bond interactions between protons belonging to the NH backbone groups or to the NH₂ groups and the phosphate surface O atoms are also established. Geometrical details are shown in Figure 3 and 4.

Figures 5 and 6 show the structures of the H-P1 and H-P2 peptides when adsorbed on the HA (001) and (010) surfaces. As anticipated (*vide supra*), the peptide groups interacting with the surfaces belong to the K and E amino acidic residues, *i.e.*, NH₂ and COOH, respectively. Significantly, for the COOH groups belonging to the E residues, spontaneous proton transfer towards the surface PO₄ groups occurs, leading to the formation of strong H-bond interactions between the protonated phosphate and the resulting COO⁻ groups. Proton transfer processes have already been predicted for glycine and glutamic acid interacting with HA surfaces.^{35,45,47,48} It is worth noting that the proton transfer process would have not been predicted by using classical force field as bond making/breaking is not taken into account. The proton transfer has important

effects on the interaction energy as, for the helix cases, an extra charge-charge component to the interaction between the negative peptide and the positive HA surface takes place. Indeed, by using Mulliken charges, it turned out that the H-P1 exhibits a net charge of $-0.8450 e$ due to the proton transfer towards the HA(001) surface (which becomes positively charged by the same amount to ensure electro neutrality). For the adsorption of the H-P2 case, the net charge ($-1.800 e$) is almost doubled with respect to the H-P1 peptide due to the presence of two E groups releasing their protons to the surface. For the HA(010) surface the above net charge values are higher (in absolute term), namely -1.3666 and -2.5103 electrons. The reason for the charge transfer enhancement when passing from HA(001) to HA(010) is revealed by inspection of the geometries of Figure 5 and 6, respectively. For the HA(001) the NH_2 group of the K residue only interacts through the N lone pair, releasing electronic charge towards the surface (inset K1 and K2, Figure 5). For the HA(010), on the contrary, a strong H-bonding is established between the NH bond and the surface basic oxygen which reverse the flux of charge (inset K1, Figure 6) and account for the higher charge transfer. Interestingly, for the random coil RC-P1/RC-P2 cases the net charges are far smaller (in absolute term): $-0.1141/-0.0047 e$ and $-0.4281/-0.5918 e$ for the HA(001) and HA(010) surfaces, respectively. Irrespective on the folding conformation, the number of interactions involving Ca^{2+} ions is higher for HA (010) surface compared with the (001) one, as the former is known to be more reactive.^{35,50} One delicate point is whether the helix structure is disrupted by the strong interactions with the HA surfaces. As Figure 5 and 6 show, the helix conformation is highly conserved upon interaction with the HA surfaces, which is also highlighted by the Ramachandran plot (Figure 2l and 2m, respectively).

The energetic features of the peptides adsorption have been collected in Table 2. The $\Delta E_{\text{ads}}^{\text{C}} + \Delta D$ values are all large and negative, *i.e.* the peptide adsorption on both HA surfaces are highly exoenergetic. The most favorable ones are obtained on the HA (010) surface, with values almost doubled with respect to the HA (001) surface, in agreement with the higher reactivity of the former compared to the latter.^{35,45} An important trend in the computed adsorption energy is shown by data of Table 2: both RC-P1 and RC-P2 exhibit close $\Delta E_{\text{ads}}^{\text{C}} + \Delta D$ values, *i.e.*, they do not depend much on the kind of mutant. The similarity is striking for the HA (001) case and less so for the HA (010) one. In contrast, H-P1 and H-P2 show large differences in the adsorption energy, with the most negative values for the H-P2 case, irrespective on the considered HA surface. The reasons are simple: for the random coil cases, the interactions are all very similar, occurring between the Ca^{2+} ions and C=O carbonyl groups of the peptides backbone. For the helix cases, on the contrary, because the interactions are through the E and K residues, it is obviously more favorable for H-P2 exhibiting four residues compared to only two for H-P1 (see Figure 5 and 6). Dispersive contribution to the interactions (ΔD , Table 2) is also remarkable, particularly for the RC peptides in comparison with the H ones. For the former, the ΔD values are more negative due to the closeness of the random coil peptides to the HA surfaces compared to the helix structures, with the exception of H-P2 on the HA (010) surface (see Figure 5 and 6). Despite the adoption of a rather flexible polarized triple zeta basis set the BSSE is still rather large and has been accounted for to improve the accuracy of the adsorption energy.

As it has been shown in the previous section, the zwitterionic form of the free P1 and P2 peptides is unstable with respect to the neutral form. An interesting point is to establish whether the adsorption may stabilize the zwitterionic form. Results are shown in Table S1 of the SI, limited to the adsorption of the helix conformers only. It turned

out that the interaction energies are almost identical for the H-P1 and in favor of the neutral form for the H-P2 case. The corresponding structures are shown in Figure S3 of the SI. The mechanism of interaction is different with respect to the neutral case, as the NH_3^+ establishes three H-bonds with the basic oxygen of the surface while the COO^- bridges two Ca^{2+} surface ions.

Structures and energetics of peptides adsorbed on micro-solvated HA surfaces

As anticipated, the present study cannot be extended by including a large number of water molecules to mimic the role of solvation on the considered processes, due to the exceedingly high computational cost. A simple algorithm based on distance criteria has been used to compute the number of H_2O molecules that would be needed to dress the adsorbed peptides up to a second solvation shell, resulting in about 100 H_2O molecules which is outside the capability of the present approach. A further complication to handle a high number of water molecules is the change in the canonical state of the amino acids with a probable stabilization of zwitterions originating by proton transfer from Glu towards Lys amino acids. Nonetheless, to at least account for the relevance of water interaction on the most active sites, the adsorption of P1 and P2 has been studied in a micro-solvation regime, *i.e.* in presence of 8 H_2O molecules pre-adsorbed at the HA (001) surface only. Indeed, for the HA (010) surface, water spontaneously dissociates,⁵¹ which would render the calculations rather complex due to important reconstruction of the pristine (010) surface upon H_2O adsorption. The 8 H_2O molecules were adsorbed in pairs on the most exposed Ca^{2+} ions of the HA(001) surface (Figure S4, SI), reducing the Lewis acidity of these sites compared to the pristine surface. The optimized structures are shown in Figures 7 and 8. For the random coil peptides (Figure 7), the interactions are similar to those occurring in dry conditions

(*vide supra*), with the $\text{Ca}^{2+}\cdots\text{O}=\text{C}$ interactions still existing despite the water molecules directly interacting with the same Ca^{2+} ion: the only change was a longer Ca-O bond distances compared to the adsorption on dry conditions. For the HW/RC-P1 case (HW=HA(001)/8W, wet surface), the presence of water induced the formation of a new $\text{Ca}^{2+}\cdots\text{NH}$ bond by establishing a H-bond with a nearby NH peptide group bringing the N atom close enough (2.650 Å) to a nearby surface Ca^{2+} ion (see region A, Figure 7).

The helix/HA surface structures are also very similar to those resulting by adsorption on dry surfaces as the binding through the K and E residues still involved Ca-N and Ca-O electrostatic interactions. Noticeably, proton transfers from the COOH groups still occurred followed by H-bonding between the surface proton and the O atom of the carboxylate moiety. As for the adsorption on dry surfaces, the P1 and P2 mutants still maintain their helix conformation as shown by Figure 8 and by the Ramachandran plot (Figure 2n).

Table 3 reports the energetic features for the adsorption process of P1 and P2 to the wet HA (001) surface. The adsorption energies are all negative showing again the exoenergetic character of the adsorption processes. Charge transfer is also very similar to that computed for the dry HA(001) surface for both random coil and helix conformations (see Table S2 of the SI). Comparison with data of Table 2 for the adsorption on dry surfaces reveals a rather subtle effect played by the adsorbed H_2O on differentiating between random coil and helix peptides. Indeed, the interaction energies are more negative for the random coil structures for the wet surfaces compared to the dry ones, whereas the reverse resulted for the helix case. This was caused by the energetic balance between the cost of displacing water from the Ca^{2+} ions and the gain due to the new H-bond interactions established by H_2O with the adsorbed peptide.

The next step towards a more realistic situation was to add four H₂O molecules to micro-solvate the interaction sites of the random coil and helix peptides (optimized structures in Figure S5, SI). To simplify the calculations, we computed the new interaction energies considering as reactants, the solvated peptide *plus* the solvated HA (001) surface and, as products, the former peptide adsorbed on the solvated HA (001) surface *plus* four free H₂O molecules derived from de-solvating the peptide and being incorporated in the surrounding water. In this way, data on the lowest half of Table 3, include the cost of de-solvating the peptide induced by the surface interaction. The interaction energies are still all negative (exoenergetic reactions) but the absolute values are all much smaller than the data not envisaging peptide solvation (data on the highest half of Table 3) and those for the adsorption on dry surfaces (see Table 2). It is worth noting that the main order of stability between the considered processes is the same for the most realistic process (data of Table 3) compared with reactions carried out in dry conditions (data of Table 2), supporting the validity of the latter approach.

Data from Table 2 and 3 can be combined in such a way to highlight the relative affinity towards the HA(001) surface of the random coil *vs* the helix conformation. Table 4 shows that the adsorption of the random coil RC-P1 on both dry and wet HA (001) and HA (010) surfaces is preferred over the H-P1 by a significant amount. The situation changes when the P2 is involved: in all cases the H-P2 is preferred over the random coil case. These trends are the same also when free energies are computed by correcting the electronic energies with the standard statistical mechanics formulae based on the adoption of harmonic frequencies computed for the free peptides in the two forms. The main message is that P2 can be stabilized as a helix by the HA surface over its most stable random coil form, at variance with the case of P1. The reasons are due to a higher number of “active sites” present in P2 with respect to P1, which can interact

more strongly *via* a subtle interplay between dispersive, electrostatic and H-bond interactions on the HA surface sites. This conclusion remains effective also when water solvation is accounted for by means of a limited number of explicit water molecules.

Concluding Remarks

The capability of HA (001) and (010) surfaces to adsorb a peptide model and drive its conformation from the most stable random coil in gas-phase to a helix due to adsorption has been studied by means of quantum mechanical B3LYP calculations carried out on periodic models of the HA surfaces. Two short peptides (P1 and P2) made by 12 amino acidic residues (P1: C-tGGKGGGGGGEGGN-t; P2: C-tGGKGGKEGGEGGN-t) have been designed *in silico* for this purpose and their preferred conformation in gas-phase has been established first, by classical molecular dynamics simulation and then, by full quantum mechanical B3LYP optimization. Exhaustive search revealed that the preferred conformation for both peptides was a random coil one, the helix ones being $\approx 100 \text{ kJ mol}^{-1}$ higher in energy. The main reasons for the random coil stability are the higher number of intra-molecular H-bonds and more effective dispersive interactions due to its compactness compared to the helix. Adsorption of P1 and P2 peptides on the HA (001) and (010) surfaces is exoenergetic for both random coil and helix conformations and for both surfaces, the (010) one showing more negative interaction energies than the (001) one. Addition of 8 H₂O molecules on the most active Ca²⁺ sites (two H₂O *per* surface Ca²⁺) at the HA (001) surface does not hinder the adsorption process, which remains exoenergetic for all cases. Despite the reduced Lewis acidity of the hydrated Ca²⁺ ions, the peptides are still able to interact with the Ca²⁺ ions by displacing the H₂O molecules, these latter engaging new H-bonds with the adsorbed proteins. For the peptides adsorbed as random

coil, the interactions are between C=O backbone groups and surface Ca^{2+} ions with large and favorable dispersive interactions of similar strength. The situation is entirely different when the peptides are adsorbed as helix, since the important interactions are through the K and E amino acidic residues (*i.e.* establishing two and four contact points for P1 and P2, respectively) and dispersion forces, which are more relevant for H-P2 than for H-P1. The charge transfer is very large for the adsorption of P1 and P2 as helix on both surfaces either dry or wet and less so when adsorbed as random coil. The charge transfer is dominated by the occurrence of the proton transfer and by the strength of the H-bonds between peptides and the HA surfaces which would have been missed by the adoption of classical force fields. To come close to more realistic conditions, also the peptides have been micro-solvated using four H_2O molecules which implied a further de-solvation cost after adsorption at the HA surfaces. All in all, the balance between the energetic cost of passing from random coil to helix, to displace the pre-adsorbed H_2O at the HA surface together with the protein desolvation and the energy gain due to the protein/surface interactions is in favor of adsorption in all cases, with the $\text{HW} + \text{RC-P1}/4\text{W} \rightarrow \text{HW}/\text{H-P1} + 4\text{W}$ process being at the limit of exoenergetic. When equilibrium between adsorbed random coil and helix peptides are established, it turned out that P1 prefers to stay adsorbed as a random coil, at variance with P2 for which the adsorption on HA (001) surface favors the helix conformation.

In conclusion, for the helix stabilization to occur over the “native” random coil one, the peptide should contain enough amino acidic residues with the proper acid/basic character and to be oriented in a favorable way to maximize the interaction with the HA surfaces. Notwithstanding, it is relevant also to address a number of important weaknesses and assumptions made in the present work due to the limits imposed by the high cost of the adopted B3LYP methodology and of the intrinsic difficulties associated

to this problem. First, the charge screening effects due to adsorbed water at the HA surfaces may decrease the favorable interactions of the helix over the random coil conformer and, at the same time, strongly polarize the water molecules directly attached to the surface cations, as shown in a recent work dealing with peptide adsorbed on oxidized silicon and amorphous silica surfaces.⁵² In our approach, two H₂O molecules per Ca atom have been considered, which only accounts for a decreased Lewis acidity of each surface cation, the amino acid residues still being able to directly contact the HA Ca²⁺ ions. The question is whether an adsorbed H₂O monolayer might completely exclude the direct contact between surface cations and the amino acid residues. We are not inclined to that possibility as we have shown recently⁴⁶ that even glycine (the simplest AA) does prefer to displace water from the HA surface to directly interact with the surface sites rather than being adsorbed on top of a layer of pre-adsorbed water molecules. Furthermore, recent work using ReaxFF force field by Monti and coworkers²⁸ shows that during long MD simulations of a fully water solvated glycine and di-glycine adsorbed on a TiO₂ surface, water does not displace the amino acids interacting directly with the surface. Second, the entropic contribution to adsorption free energy due to water displacement upon binding has not been considered. As a rough guess, the contact surface area between the peptide and the HA surface is a measure of the amount of displaced water. This quantity is higher for the RC than for the H peptide, which means that the adsorption of the former is entropically favoured over the latter one. To what extent this may change the present results is however difficult to assess quantitatively. Third, the role of solvent on the folding of free P1 and P2 peptide has not been accounted for here. As expected for such a small peptides, classical MD calculations (Figure S6, SI) revealed a different behavior of the considered peptides in gas-phase and in water. That is, a different pool of conformations appeared as a function

of the simulation medium, but, significantly, all conformations exhibiting a random coil character while helix structures were never present, neither in gas-phase nor in solution. As we are interested to see which key interactions are stabilizing the helix over a random coil (whatever it will be) the main message of this work is therefore preserved. In conclusion, the present results, despite various compromises needed to run large quantum mechanical calculations which prevent the adoption of fully solvated systems, are in line with the experimental results of Capriotti et al.,¹⁵ where the stabilization of the α -helix conformation induced by HA surfaces of a de novo design peptide rich in γ -carboxyglutamic acid residues, was observed.

It is worth mentioning that this work addresses the specific case of HA surfaces as helix-stabilizing substrates; however, other surfaces capable to establish additional interactions that compensate the folding energy costs are also suitable platforms for stabilizing folded conformations. This is for instance the case of silica nanoparticles; *i.e.*, deprotonated surfaces (with Si-O⁻ terminations) are able to stabilize helix conformations of arginine- and lysine-rich peptides (in solution these residues are protonated) due to the strong electrostatic interactions between the partners.^{11,12}

Acknowledgements

AR is indebted to MICINN for a postdoctoral Juan de la Cierva contract and for the CTQ2011-24847/BQU project. PU kindly acknowledges BSC-MN for the generous allowance of computing time through the “BCV-2008-2-0013: Simulation of peptide folding induced by inorganic materials” and “BCV-2008-2-0013: Simulation of peptide folding induced by inorganic materials (II)” projects.

Supporting Information Available

Computational details, supplementary structures, absolute B3LYP energies and coordinates of all considered structures, are provided in the Supplementary Information (SI). This information is available free of charge via the internet at <http://pubs.acs.org>.

Table 1. Reaction energies (electronic ΔE_{rx} , enthalpies ΔH_{rx} , and Gibbs ΔG_{rx}) for the conversion of peptides from their random coil structures to helix forms in gas-phase. Data computed at B3LYP/VTZP on structures optimized at B3LYP/double- ζ polarized level. Values in parenthesis are for fully B3LYP/VTZP optimized structures. Units in kJ mol^{-1} .

Folding process	ΔE_{rx}	ΔD	$\Delta E_{rx} + \Delta D$	$\Delta H_{rx} + \Delta D$ ^a	$\Delta G_{rx} + \Delta D$ ^a
RC-P1 \rightarrow H-P1	45.1	48.9	94.0	78.3	101.7
	(49.6)	(48.9)	(98.5)	(82.8)	(106.2)
RC-P2 \rightarrow H-P2	52.1	31.0	83.1	72.4	93.6
	(56.0)	(31.0)	(87.0)	(76.3)	(97.5)

^a Values computed by adding to the $\Delta E_{rx} + \Delta D$ values the enthalpic and entropic corrections obtained from frequency calculations performed at B3LYP/double- ζ level and T= 298 K.

Table 2. Adsorption of dry P1 and P2 peptides on dry HA(001) and HA(010) hydroxyapatite surfaces (H). ΔE_{ads} , and $\Delta E_{\text{ads}}^{\text{C}}$ as non-corrected and BSSE corrected interaction energies, respectively. RC-P1, RC-P2 and H-P1 and H-P2 label the random coil and helix peptide conformations, respectively. %BSSE is the adopted BSSE percentage to the ΔE_{ads} , ΔD is the dispersion contribution and $\Delta E_{\text{ads}+\Delta D}^{\text{C}}$ the final adsorption energies inclusive of dispersion. Units in kJ mol^{-1} .

Adsorption process	ΔE_{ads}	$\Delta E_{\text{ads}}^{\text{C}}$	%BSSE	ΔD	$\Delta E_{\text{ads}+\Delta D}^{\text{C}}$
HA(001) dry surface					
H + RC-P1 \rightarrow H/RC-P1	-179.1	-93.6	48	-144.1	-237.7
H + RC-P2 \rightarrow H/RC-P2	-232.9	-144.2	38	-101.5	-245.7
H + RC-P1 \rightarrow H/H-P1	-306.3	-159.3 ^a	48	-19.9	-179.2
H + RC-P2 \rightarrow H/H-P2	-530.5	-275.9 ^a	48	-85.8	-361.7
HA(010) dry surface					
H + RC-P1 \rightarrow H/RC-P1	-499.1	-281.8	43	-210.5	-492.3
H + RC-P2 \rightarrow H/RC-P2	-523.9	-308.8	41	-237.7	-546.5
H + RC-P1 \rightarrow H/H-P1	-546.4	-311.4 ^a	43	-145.7	-457.1
H + RC-P2 \rightarrow H/H-P2	-911.2	-519.4 ^a	43	-240.5	-759.9

^a Estimated $\Delta E_{\text{ads}}^{\text{C}}$ values obtained by assuming the highest BSSE percentage computed for the RC systems. The actual $\Delta E_{\text{ads}}^{\text{C}}$ values cannot be computed due to the deprotonation of the helix upon HA adsorption (see methods section for more details).

Table 3. Adsorption of dry and wet (4 H₂O per peptide) P1 and P2 peptides on micro-solvated (8 H₂O per unit cell) HA(001) hydroxyapatite surfaces (HW). ΔE_{ads} , and $\Delta E_{\text{ads}}^{\text{C}}$ as non-corrected and BSSE corrected interaction energies, respectively. RC-P1, RC-P2, H-P1 and H-P2 label the random coil and helix peptide conformations, respectively. %BSSE is the adopted BSSE percentage to the ΔE_{ads} , ΔD is the dispersion contribution and $\Delta E_{\text{ads}}^{\text{C}} + \Delta D$ the final adsorption energies inclusive of dispersion. Units in kJ mol⁻¹.

Adsorption process	ΔE_{ads}	$\Delta E_{\text{ads}}^{\text{C}}$	%BSSE	ΔD	$\Delta E_{\text{ads}}^{\text{C}} + \Delta D$
wet HA(001) surface/dry peptide					
HW + RC-P1 → HW/RC-P1	-202.4	-128.4	37	-171.4	-299.8
HW + RC-P2 → HW/RC-P2	-195.9	-139.3	29	-169.1	-308.4
HW + RC-P1 → HW/H-P1	-200.5	-126.3 ^{a,b}	37	-20.4	-146.7
HW + RC-P2 → HW/H-P2	-371.5	-234.0 ^{a,b}	37	-121.4	-355.4
wet HA(001) surface/wet peptide					
HW + RC-P1/4W → HW/RC-P1 + 4W	-31.4	-19.8 ^{a,c}	37	-134.9	-154.7
HW + RC-P2/4W → HW/RC-P2 + 4W	-15.4	-9.7 ^{a,c}	37	-122.3	-132.0
HW + RC-P1/4W → HW/H-P1 + 4W	-29.5	-18.6 ^{a,c}	37	+16.1	-2.5
HW + RC-P2/4W → HW/H-P2 + 4W	-191.0	-120.3 ^{a,c}	37	-74.6	-194.9

^a Estimated $\Delta E_{\text{ads}}^{\text{C}}$ values obtained by assuming the highest percentage of BSSE computed for the RC systems.

^b The actual $\Delta E_{\text{ads}}^{\text{C}}$ values cannot be computed due to the de-protonation of the helix upon HA adsorption (see methods section for more details).

^c The actual $\Delta E_{\text{ads}}^{\text{C}}$ values cannot be computed due to the expulsion of 4 water molecules.

Table 4. Folding energies (electronic ΔE_{rx} , enthalpies ΔH_{rx} , and Gibbs ΔG_{rx}) for the P1 and P2 peptides induced by the adsorption at the dry HA (001) and (010) surfaces (H) and on the wet HA (001) one (HW=HA(001)/8W). RC-P1, RC-P2, H-P1 and H-P2 label the random coil and helix peptide conformations, respectively
Units in kJ mol^{-1} .

Folding process	ΔE_{rx}^a	ΔH_{rx}^b	ΔG_{rx}^c
HA(001) dry surface			
H/RC-P1 \rightarrow H/H-P1	+58.5	+42.8	+66.1
H/RC-P2 \rightarrow H/H-P2	-116.0	-126.7	-105.5
HA(010) dry surface			
H/RC-P1 \rightarrow H/H-P1	+35.2	+19.5	+42.8
H/RC-P2 \rightarrow H/H-P2	-213.4	-224.1	-202.9
HA(001)/W8 wet surface			
HW/RC-P1 \rightarrow HW/H-P1	+152.2	+136.5	+159.8
HW/RC-P2 \rightarrow HW/H-P2	-62.9	-73.6	-52.4

^a From the corresponding $\Delta E_{ads}^C + \Delta D$ values.

^b From the corresponding $\Delta E_{ads}^C + \Delta D$ values, including the enthalpic corrections obtained from the free-surface processes (Table 1).

^c From the corresponding $\Delta E_{ads}^C + \Delta D$ values, including the entropic corrections obtained from the free-surface processes (Table 1).

Captions to the Figures

Figure 1. Scheme for the *in silico* design of the helix structures. H-[G]_p: periodic helix polyglycine, H-[G]_m: a finite 12-mer glycine α -helix (terminus labeled by star character), H-P1: mutant with one glutamic acid and one lysine residues, H-P2: mutant with two glutamic acid and two lysine residues. For a proper interaction with the HA surfaces the residues are oriented towards the same side of the main helix axis.

Figure 2. Top: (a) and (b): H-P1 and H-P2 optimized helix structures viewed normal to main helix axis. (c)/(d) and (e)/(f): same as sections (a) and (b) viewed along the main helix axis. Region nearby the N-t and C-t adopt a 3_{10} and α -helix structures, respectively. (g) and (h): skeleton view of the P1 and P2 optimized structures in their “native” random coil conformations. Bottom: Ramachandran plots of gas-phase P1 and P2 peptides in their helix H-P1 (empty diamond), H-P2 (empty circle) and random coil RC-P1 (filled square), RC-P2 (filled triangle) conformations as: (i) free gas-phase; (l) adsorbed on HA (001) surface; (m) adsorbed on HA (010) surface; (n) adsorbed on wet HA(001)/W8 surface (two H₂O per Ca²⁺).

Figure 3. B3LYP optimized structures of RC-P1 and RC-P2 adsorbed on the (001) hydroxyapatite surface. Insets A, B and C highlight the shortest intermolecular contacts between peptides and the surface atoms. Bottom: backbone only view of the adsorbed peptides.

Figure 4. B3LYP optimized structures of RC-P1 and RC-P2 adsorbed on the (010) hydroxyapatite surface. Insets A, B, C, D and E highlight the shortest intermolecular

contacts between peptides and the surface atoms. Bottom: backbone only view of the adsorbed peptides.

Figure 5. B3LYP-optimized structures of H-P1 and H-P2 adsorbed on the (001) hydroxyapatite surface. Insets K1, K2, E1 and E2 highlight the shortest intermolecular contacts between peptides and the surface atoms. Bottom: backbone only view of the adsorbed helix peptides.

Figure 6. B3LYP optimized structures of H-P1 and H-P2 adsorbed on the (010) hydroxyapatite surface. Insets K1, K2, E1 and E2 highlight the shortest intermolecular contacts between peptides and the surface atoms. Bottom: backbone only view of the adsorbed helix peptides.

Figure 7. B3LYP-optimized structures of RC-P1 and RC-P2 adsorbed on the wet (001) hydroxyapatite surface (8 H₂O molecules). Water molecules labeled by W and color coded in cyan. Insets A, B, C and D highlight the shortest intermolecular contacts between peptides and the surface atoms. Bottom: backbone only view of the random coil adsorbed peptides.

Figure 8. B3LYP optimized structures of H-P1 and H-P2 adsorbed on the (001) hydroxyapatite surface (8 H₂O molecules). Water molecules labeled by W and color coded in cyan. Bottom: backbone only view of the adsorbed helix peptides.

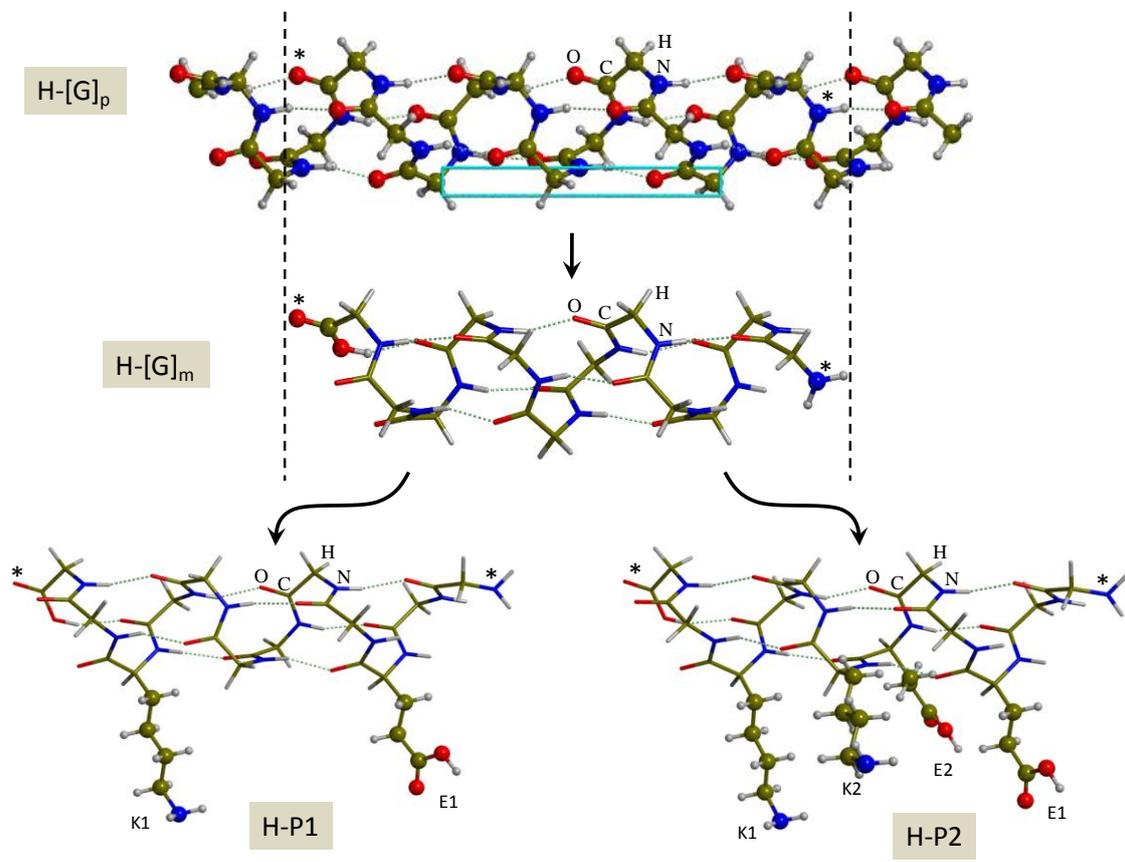


Figure 1

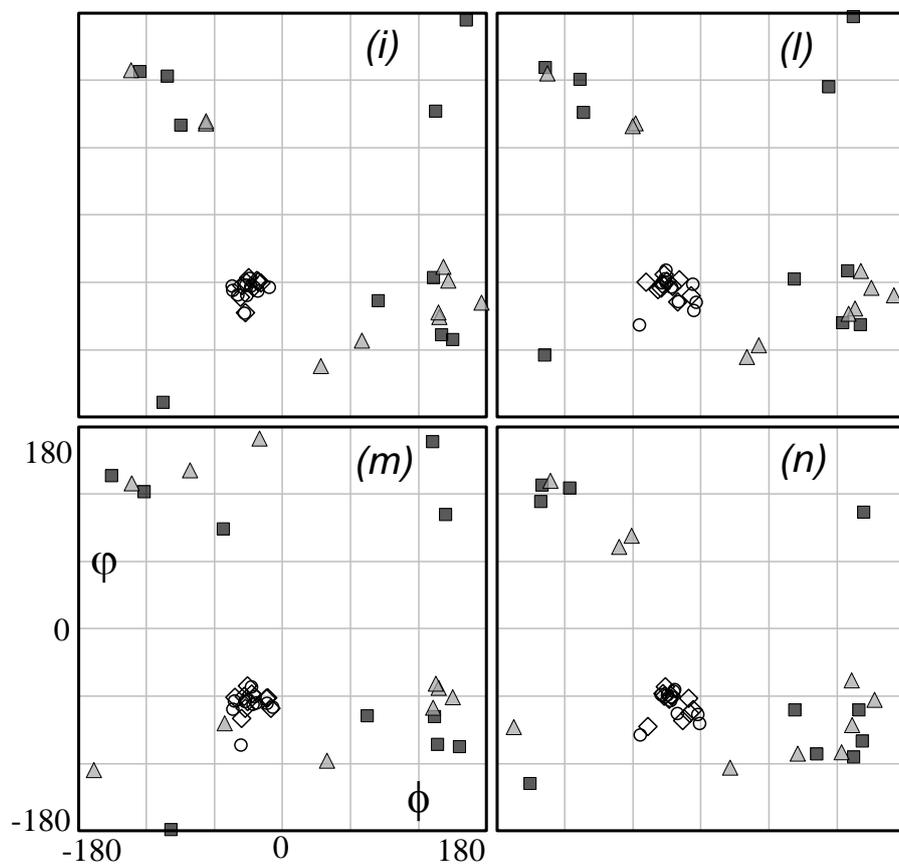
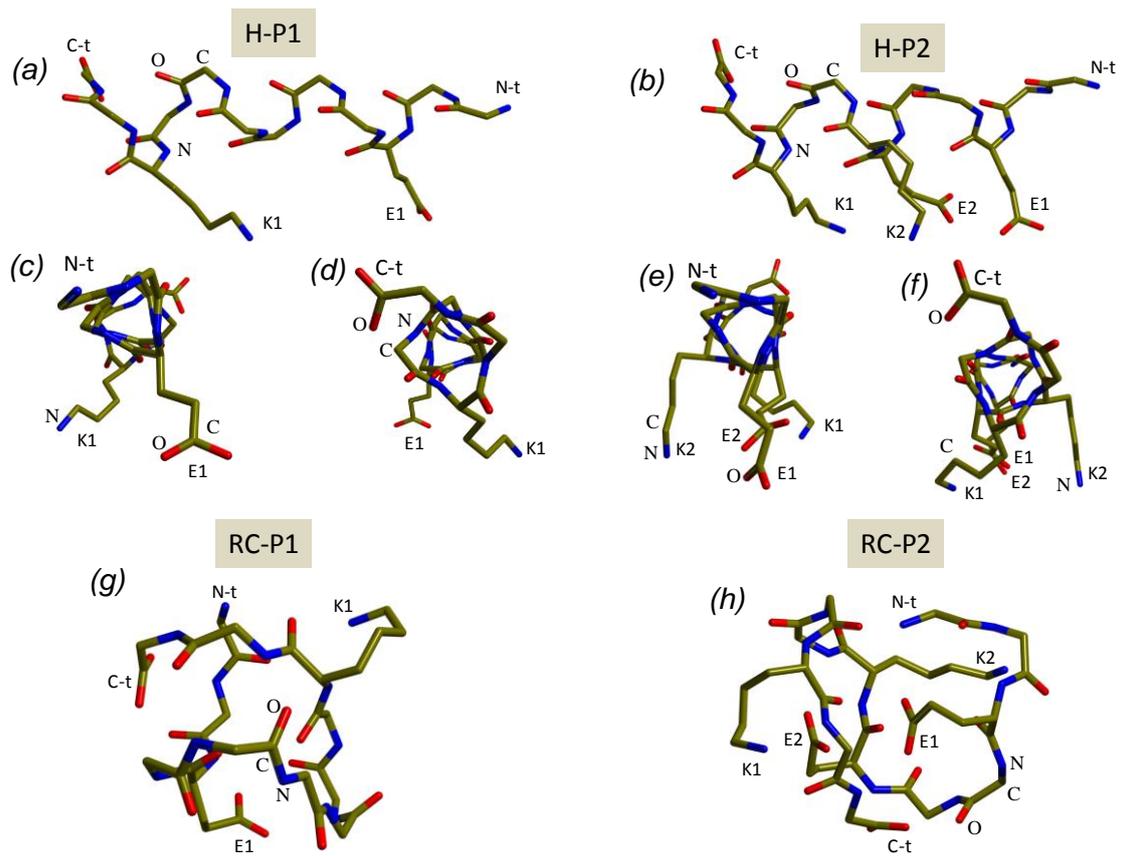
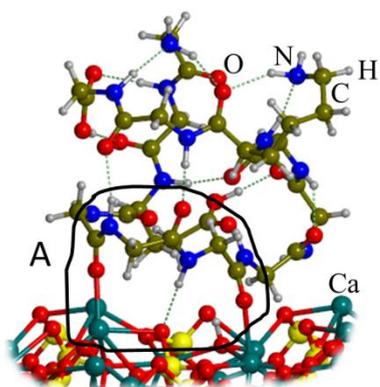


Figure 2

HA001/RC-P1



HA001/RC-P2

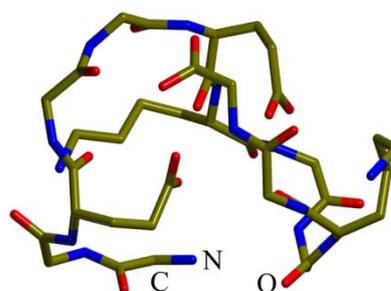
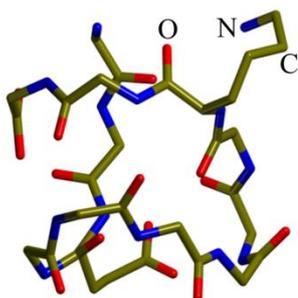
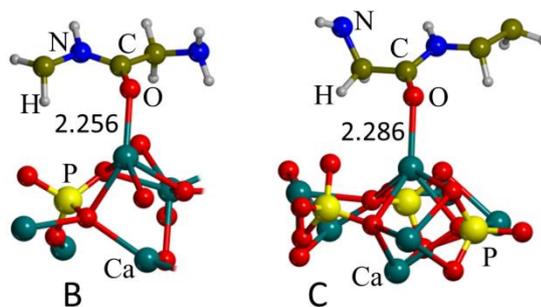
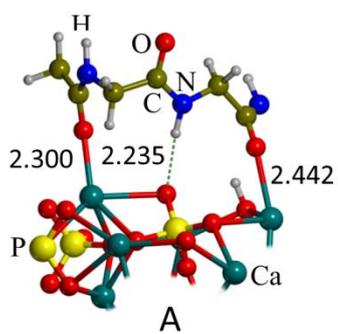
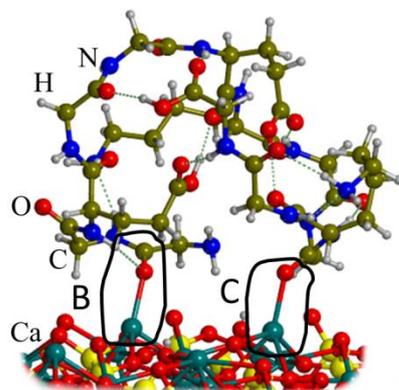


Figure 3

HA010/RC-P1

HA010/RC-P2

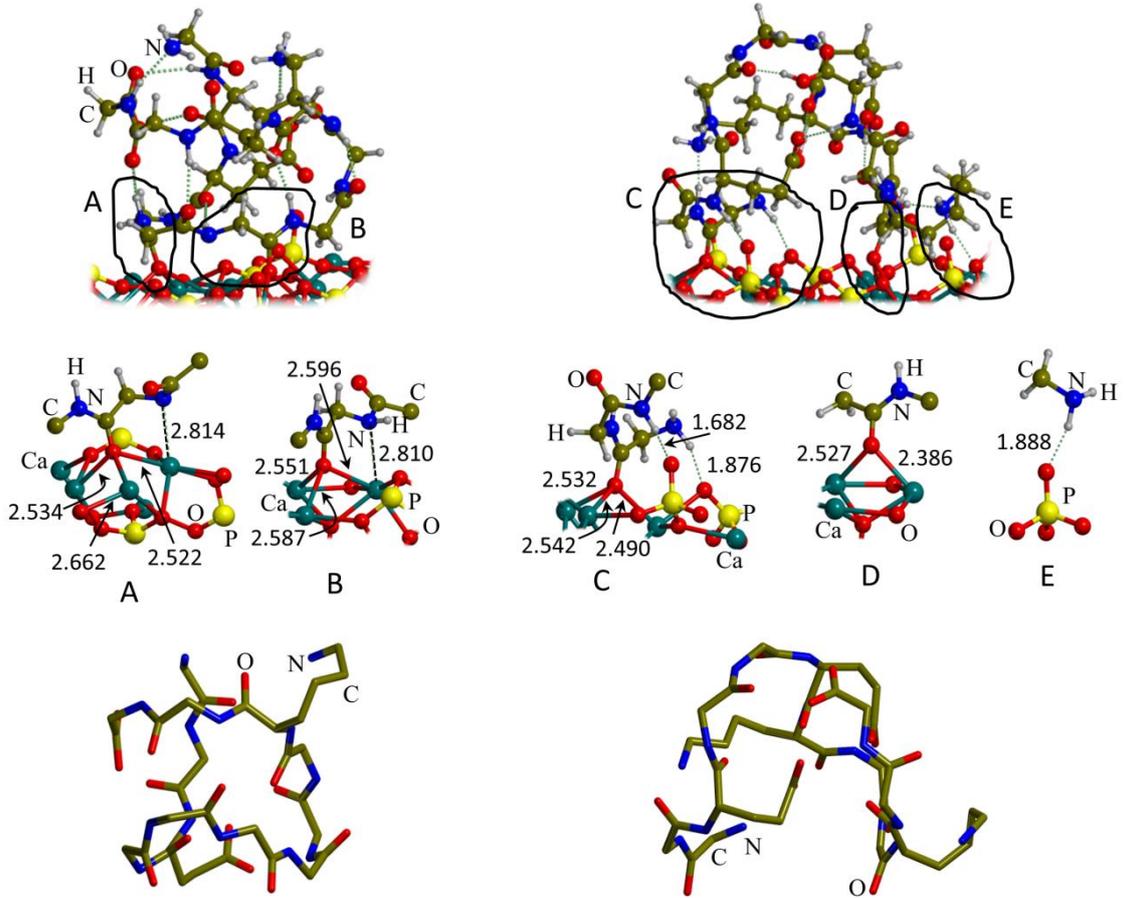


Figure 4

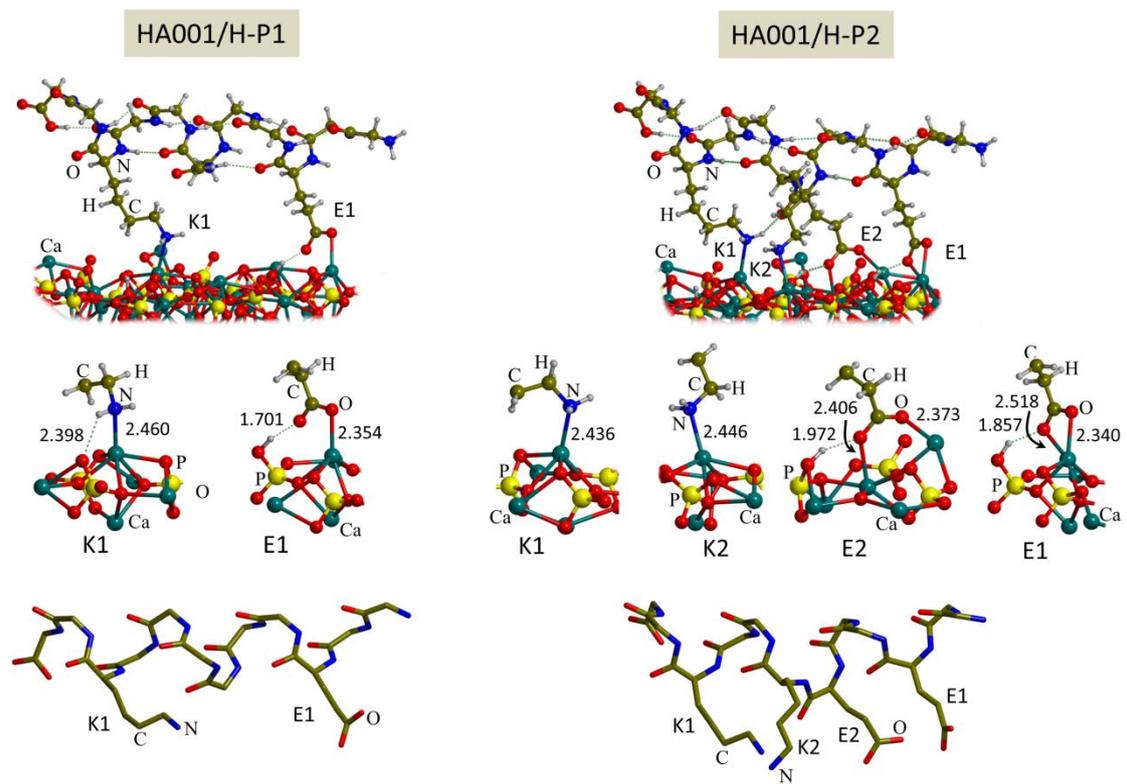


Figure 5

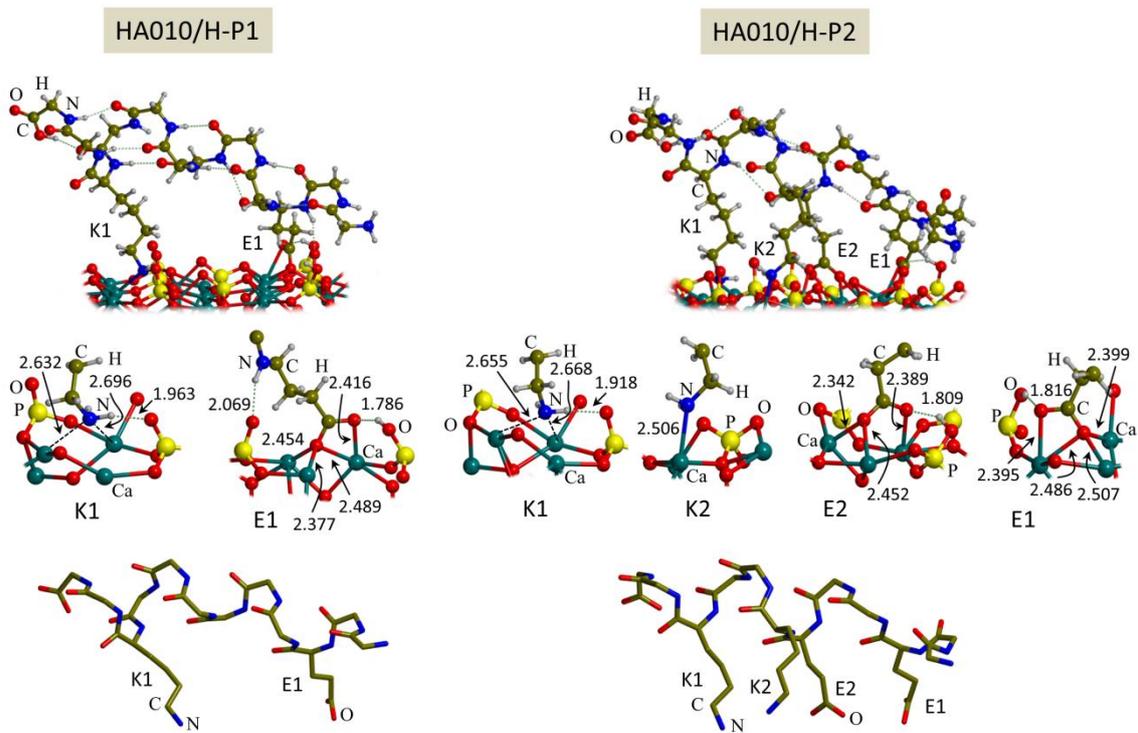
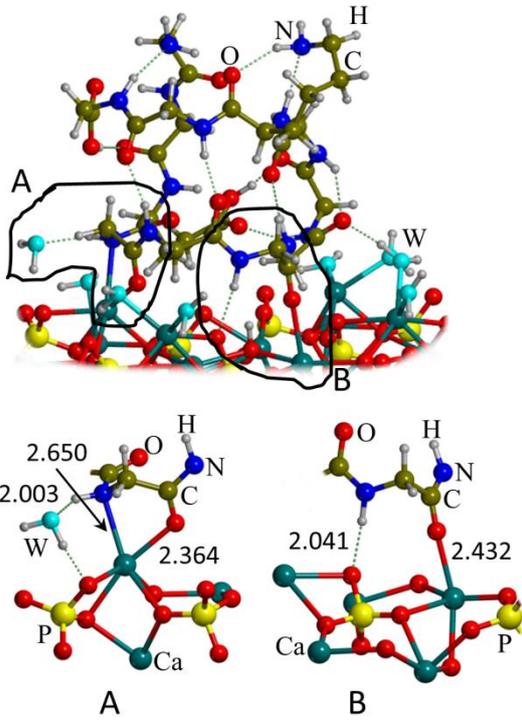


Figure 6

HA001/8W/RC-P1



HA001/8W/RC-P2

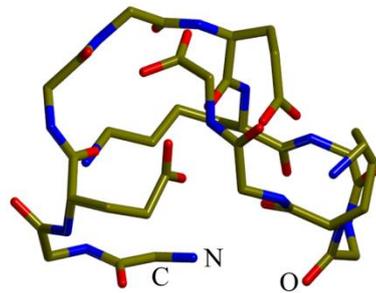
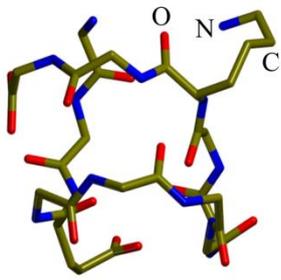
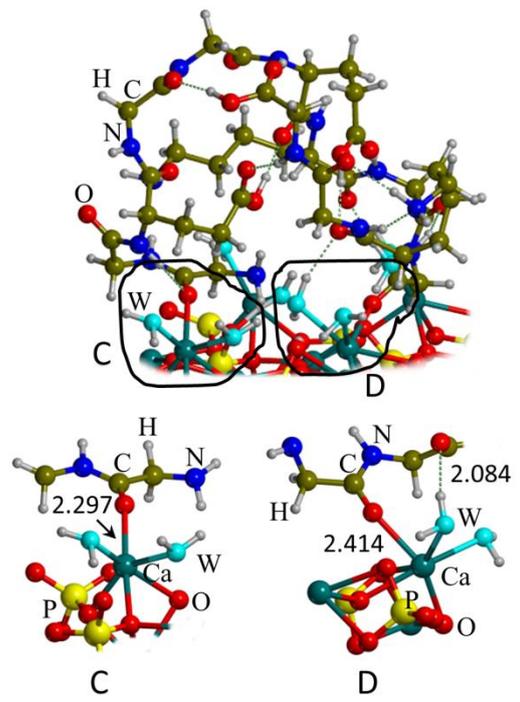


Figure 7

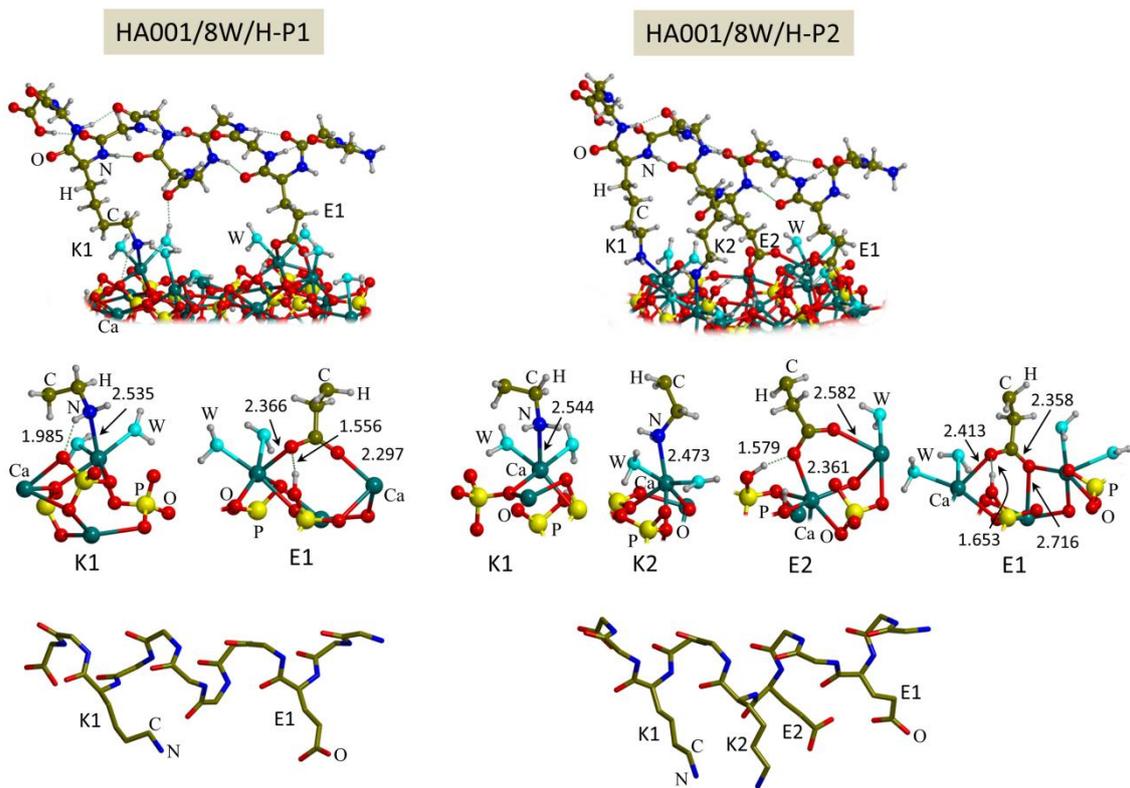


Figure 8

References

- (1) Cline, D. J.; Thorpe, C.; Schneider, J. P. *J. Am. Chem. Soc.* **2003**, *125*, 2923.
- (2) Cox, E. H.; McLendon, G. L. *Curr. Opin. Chem. Biol.* **2000**, *4*, 162.
- (3) Kelso, M. J.; Hoang, H. N.; Oliver, W.; Sokolenko, N.; March, D. R.; Appleton, T. G.; Fairlie, D. P. *Angew. Chem. Int. Ed.* **2003**, *42*, 421.
- (4) Lombardi, A.; Marasco, D.; Maglio, O.; Di Costanzo, L.; Natri, F.; Pavone, V. *Proc. Natl. Acad. Sci. USA* **2000**, *97*, 11922.
- (5) Ojida, A.; Inoue, M.-a.; Mito-oka, Y.; Hamachi, I. *J. Am. Chem. Soc.* **2003**, *125*, 10184.
- (6) Nicoll, A. J.; Miller, D. J.; Fütterer, K.; Ravelli, R.; Allemann, R. K. *J. Am. Chem. Soc.* **2006**, *128*, 9187.
- (7) Le Clainche, L.; Plancque, G.; Amekraz, B.; Moulin, C.; Pradines-Lecomte, C.; Peltier, G.; Vita, C. *J. Biol. Inorg. Chem.* **2003**, *8*, 334.
- (8) Jackson, D. Y.; King, D. S.; Chmielewski, J.; Singh, S.; Schultz, P. G. *J. Am. Chem. Soc.* **1991**, *113*, 9391.
- (9) Long, J. R.; Oyler, N.; Drobny, G. P.; Stayton, P. S. *J. Am. Chem. Soc.* **2002**, *124*, 6297.
- (10) Dolain, C.; Hatakeyama, Y.; Sawada, T.; Tashiro, S.; Fujita, M. *J. Am. Chem. Soc.* **2010**, *132*, 5564.
- (11) Lundqvist, M.; Nygren, P.; Jonsson, B.-H.; Broo, K. *Angew. Chem. Int. Ed.* **2006**, *45*, 8169.
- (12) Nygren, P.; Lundqvist, M.; Broo, K.; Jonsson, B.-H. *Nano Lett.* **2008**, *8*, 1844.
- (13) Fillon, Y.; Verma, A.; Ghosh, P.; Ernenwein, D.; Rotello, V. M.; Chmielewski, J. *J. Am. Chem. Soc.* **2007**, *129*, 6676.
- (14) Verma, A.; Nakade, H.; Simard, J. M.; Rotello, V. M. *J. Am. Chem. Soc.* **2004**, *126*, 10806.
- (15) Capriotti, L. A.; Beebe, T. P.; Schneider, J. P. *J. Am. Chem. Soc.* **2007**, *129*, 5281.

- (16) Baugh, L.; Weidner, T.; Baio, J. E.; Nguyen, P.-C. T.; Gamble, L. J.; Stayton, P. S.; Castner, D. G. *Langmuir* **2010**, *26*, 16434.
- (17) Goobes, G.; Goobes, R.; Schueler-Furman, O.; Baker, D.; Stayton, P. S.; Drobny, G. P. *Proc. Natl. Acad. Sci. USA* **2006**, *103*, 16083.
- (18) Goobes, G.; Goobes, R.; Shaw, W. J.; Gibson, J. M.; Long, J. R.; Raghunathan, V.; Schueler-Furman, O.; Popham, J. M.; Baker, D.; Campbell, C. T.; Stayton, P. S.; Drobny, G. P. *Magn. Reson. Chem.* **2007**, *45*, S32.
- (19) Masica, D. L.; Gray, J. J.; Shaw, W. J. *J. Phys. Chem. C* **2011**, *115*, 13775.
- (20) Peczu, M. W.; Hamilton, A. D. *Chem. Rev.* **2000**, *100*, 2479.
- (21) Wächtershäuser, G. *Proc. Natl. Acad. Sci. USA* **1990**, *87*, 200.
- (22) Cody, G. D.; Boctor, N. Z.; Filley, T. R.; Hazen, R. M.; Scott, J. H.; Sharma, A.; Yoder, H. S. *Science* **2000**, *289*, 1337.
- (23) Makrodimitris, K.; Masica, D. L.; Kim, E. T.; Gray, J. J. *J. Am. Chem. Soc.* **2007**, *129*, 13713.
- (24) Ghiringhelli, L. M.; Hess, B.; van der Vegt, N. F. A.; Delle Site, L. *J. Am. Chem. Soc.* **2008**, *130*, 13460.
- (25) Calzolari, A.; Cicero, G.; Cavazzoni, C.; Di Felice, R.; Catellani, A.; Corni, S. *J. Am. Chem. Soc.* **2010**, *132*, 4790.
- (26) Di Felice, R.; Corni, S. *J. Phys. Chem. Lett.* **2011**, *2*, 1510.
- (27) van Duin, A. C. T.; Dasgupta, S.; Lorant, F.; Goddard, W. A. *J. Phys. Chem. A* **2001**, *105*, 9396.
- (28) Monti, S.; van Duin, A. C. T.; Kim, S.-Y.; Barone, V. *J. Phys. Chem. C* **2012**, *116*, 5141.
- (29) Rahaman, O.; van Duin, A. C. T.; Goddard, W. A.; Doren, D. J. *J. Phys. Chem. B* **2010**, *115*, 249.
- (30) Bush, I. J.; Tomic, S.; Searle, B. G.; Mallia, G.; Bailey, C. L.; Montanari, B.; Bernasconi, L.; Carr, J. M.; Harrison, N. M. *Proc. R. Soc. A-Math. Phys. Eng. Sci.* **2011**, *467* (2131), 2112.
- (31) Becke, A. D. *J. Chem. Phys.* **1993**, *98*, 5648.
- (32) Dovesi, R.; Saunders, V. R.; Roetti, C.; Orlando, R.; Zicovich-Wilson, C. M.; Pascale, F.; Civalleri, B.; Doll, K.; Harrison, N. M.; Bush, I. J.; D'Arco, P.; Llunell, M.; CRYSTAL09, University of Torino: Torino, 2009.

- (33) Dovesi, R.; Orlando, R.; Civalleri, B.; Roetti, C.; Saunders, V. R.; Zicovich-Wilson, C. M. *Z. Kristallogr.* **2005**, *220*, 571.
- (34) Lee, C.; Yang, W.; Parr, R. G. *Phys. Rev. B* **1988**, *37*, 785.
- (35) Corno, M.; Rimola, A.; Bolis, V.; Ugliengo, P. *Phys. Chem. Chem. Phys.* **2010**, *12*, 6309.
- (36) Schafer, A.; Horn, H.; Ahlrichs, R. *J. Chem. Phys.* **1992**, *97*, 2571.
- (37) Rimola, A.; Civalleri, B.; Ugliengo, P. *Phys. Chem. Chem. Phys.* **2010**, *12*, 6357.
- (38) Grimme, S. *J. Comput. Chem.* **2006**, *27*, 1787.
- (39) Civalleri, B.; Zicovich-Wilson, C. M.; Valenzano, L.; Ugliengo, P. *CrystEngComm* **2008**, *10*, 405.
- (40) Oostenbrink, C.; Villa, A.; Mark, A. E.; van Gunsteren, W. F. *J. Comput. Chem.* **2004**, *25*, 1656.
- (41) van Der Spoel, D.; Lindahl, E.; Hess, B.; Groenhof, G.; Mark, A. E.; Berendsen, H. J. *J. Comput. Chem.* **2005**, *26*, 1701.
- (42) Berendsen, H. J. C.; Postma, J. P. M.; van Gunsteren, W. F.; DiNola, A.; Haak, J. R. *J. Chem. Phys.* **1984**, *81*, 3684.
- (43) Hess, B.; Bekker, H.; Berendsen, H. J. C.; Fraaije, J. G. E. M. *J. Comp. Chem.* **1997**, *18*, 1463.
- (44) Amadei, A.; Linssen, A. B.; Berendsen, H. J. C. *Proteins* **1993**, *17*, 412.
- (45) Rimola, A.; Corno, M.; Zicovich-Wilson, C. M.; Ugliengo, P. *J. Am. Chem. Soc.* **2008**, *130*, 16181.
- (46) Rimola, A.; Corno, M.; Zicovich-Wilson, C. M.; Ugliengo, P. *Phys. Chem. Chem. Phys.* **2009**, *11*, 9005.
- (47) Rimola, A.; Sakhno, Y.; Bertinetti, L.; Lelli, M.; Martra, G.; Ugliengo, P. *J. Phys. Chem. Lett.* **2011**, *2*, 1390.
- (48) Rimola, A.; Corno, M.; Garza, J.; Ugliengo, P. *Philos. Trans. R. Soc. A* **2012**, *370*, 1478.
- (49) Ramaekers, R.; Pajak, J.; Lambie, B.; Maes, G. *J. Chem. Phys.* **2004**, *120*, 4182.
- (50) Corno, M.; Orlando, R.; Civalleri, B.; Ugliengo, P. *Eur. J. Mineral.* **2007**, *19*, 757.
- (51) Corno, M.; Busco, C.; Bolis, V.; Tosoni, S.; Ugliengo, P. *Langmuir* **2009**, *25*, 2188.

(52) Schneider, J.; Ciacchi, L. C. *J. Am. Chem. Soc.* **2012**, *134*, 2407.

TOC Graphic

